California Drug Recall Information

Recall Name
RB Recalls MUCINEX® **Due to Undeclared Levels of Active Ingredients**

Recall Date	Product Description	Recalling Firm	Recall Reason
04/21/15	• MUCINEX® FAST-MAX® Night Time Cold & Flu • MUCINEX® FAST-MAX® Cold & Sinus • MUCINEX® FAST-MAX® Severe Congestion & Cough • MUCINEX® FAST-MAX® Cold, Flu & Sore Throat	**RB (formerly Reckitt Benckiser)** Parsippany, NJ	*The products may have the incorrect Drug Facts label on the back panel.* *This could cause the consumer to be unaware of side effects and/or risks associated with ingredients including Acetaminophen, Dextromethorphan, Guaifenesin, Phenylephrine and/or Diphenhydramine.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
N/A	Product Labels Affected Lots	Nationwide	Expiry: 5/31/2016 to 1/31/2017

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm444028.htm